

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property
Organization
International Bureau



(43) International Publication Date
12 May 2005 (12.05.2005)

PCT

(10) International Publication Number
WO 2005/041861 A2

- (51) International Patent Classification⁷: **A61K**
- (21) International Application Number:
PCT/US2004/027901
- (22) International Filing Date: 27 August 2004 (27.08.2004)
- (25) Filing Language: English
- (26) Publication Language: English
- (30) Priority Data:
60/498,347 28 August 2003 (28.08.2003) US
- (71) Applicant (for all designated States except US): PURE BIOSCIENCE, INC. [US/US]; 1725 Gillespie Way, El Cajon, CA 92020 (US).
- (72) Inventor; and
- (75) Inventor/Applicant (for US only): ARATA, Andrew, B. [US/US]; Route 29, Box 2639, Lake City, FL 32024 (US).
- (74) Agents: CAMPBELL, Cathryn et al.; McDermott Will & Emery LLP, Suite 700, 4370 La Jolla Village Drive, San Diego, CA 92122 (US).
- (81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.
- (84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).
- Published:
— without international search report and to be republished upon receipt of that report
- For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

WO 2005/041861 A2

(54) Title: ANHYDROUS SILVER DIHYDROGEN CITRATE COMPOSITIONS

(57) Abstract: Anhydrous silver dihydrogen citrate compositions comprise silver dihydrogen citrate and citric acid. The anhydrous compositions can be prepared by freeze-drying. The anhydrous compositions can be reconstituted with a suitable diluent to form silver dihydrogen citrate compositions. The anhydrous compositions can be reconstituted and applied to a variety of substrates to impart an antimicrobial effect on the substrates.

ANHYDROUS SILVER DIHYDROGEN CITRATE COMPOSITIONS

FIELD OF THE INVENTION

This invention generally relates to antimicrobial compositions. In particular, the
5 invention provides compositions comprising anhydrous silver dihydrogen citrate and
citric acid, processes of making the compositions and antimicrobial methods of using the
solid compositions.

BACKGROUND OF THE INVENTION

Antimicrobial metal ion solutions have been used as disinfectants. Silver
10 solutions have been used as disinfectants in cooling towers, swimming pools, hot water
systems in hospitals, portable water systems and spa pools. Additionally, silver ion
solutions have been prepared for the treatment of wounds, however the silver ions used in
the proposed methods are unstable and must be generated near the wound in order to
deliver a therapeutic dose to the wound site.

15 Silver salts, such as silver citrate salts, have also been proposed as antimicrobial
dusting agents. However, these dusting agents must be kept dry and are generally not
convenient for imparting preservative value to consumer products or for delivering
antimicrobial effects to an end user or to the environment of the end user. Colloidal silver
has found a variety of uses, including: as a wood preservative; as a disinfectant of food
20 and beverage containers and industrial processing equipment; as a bactericide in paints; as
a biocide in synthetic polymer films; and as a sterilizing agent in bandages.

Aqueous solutions of silver dihydrogen citrate and citric acid have been disclosed
in US Patent No. 6,197,814 (incorporated herein in its entirety) as disinfectants in a
variety of settings. These water solutions of silver dihydrogen citrate and citric acid are
25 made by passing an electrical current through a pair of silver electrodes that are immersed
in a water solution of citric acid. These silver dihydrogen citrate solutions are effective
against a wide-variety of microbes, including bacteria, viruses and fungi, and are non-
toxic in the human environment at concentrations effective to combat microbial
infestation. However, aqueous solutions are bulky and heavy, and are thus uneconomical
30 to store and to transport.

There is a need for silver dihydrogen citrate compositions that can be economically stored, transported or both. Such compositions should be substantially free of water in order to minimize the space required for their storage, as well to minimize their shipping weight. Such compositions should confer the same beneficial antimicrobial effect as solutions of silver dihydrogen citrate when combined with water or other aqueous solution. The present invention satisfies this need and provides related advantages as well.

SUMMARY OF THE INVENTION

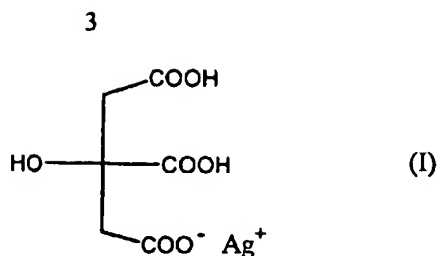
The present invention provides compositions comprising anhydrous silver dihydrogen citrate and citric acid and a process of making said compositions. The process includes freezing a stock solution of aqueous silver dihydrogen citrate and then freeze-drying the frozen stock solution to produce anhydrous silver dihydrogen citrate.

The invention further provides a method of using anhydrous silver dihydrogen citrate and citric acid by combining it with an aqueous diluent to produce an antimicrobially effective aqueous silver dihydrogen citrate solution.

DETAILED DESCRIPTION OF THE INVENTION

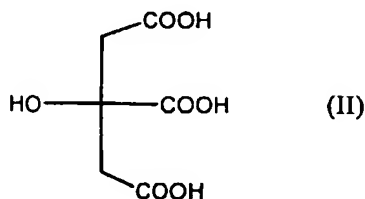
The invention provides compositions comprising anhydrous silver dihydrogen citrate and citric acid. Surprisingly, it has been found that the anhydrous silver dihydrogen citrate compositions may be reconstituted after a period of time to make stable silver dihydrogen citrate solutions having effective antimicrobial activity. The anhydrous compositions offer the advantage of ease and economy of storage. They also offer the advantage of being more economical to transport than aqueous solutions of silver dihydrogen citrate. As the anhydrous compositions may be conveniently reconstituted with water or another aqueous diluent, the anhydrous compositions offer a convenient alternative to aqueous solutions of silver dihydrogen citrate.

Silver dihydrogen citrate has been identified as the active antimicrobial ingredient in a solution produced by electrolysis of silver electrodes immersed in a citric acid electrolyte solution. The structure of silver dihydrogen citrate can be represented by formula I:



wherein Ag^+ is silver ion and the negatively charged carbonate (COO^-) is the most likely charged group in the dihydrogen citrate moiety. The chemical formula for this complex is $\text{AgC}_6\text{H}_7\text{O}_7$, and its molecular weight is 298.99 g/mol. As can be seen from the formula above, silver dihydrogen citrate is a salt, wherein the silver ion is the cation and the dihydrogen citrate ion is the anion. In solution, the salt is present in a dissociated state, the cation and anion being surrounded by water molecules, which associate with the ions through their polar OH groups.

Citric acid can be represented by formula II:



having a chemical formula of $\text{C}_6\text{H}_8\text{O}_7$ and a molecular weight of 192.13 g/mol. Citric acid is readily available from commercial sources. Citric acid is very soluble in water. Aqueous solutions of citric acid may be up to about 60% (wt/vol) citric acid in water at 25°C.

The invention also provides methods of making solid silver dihydrogen citrate compositions. The method starts with providing a stock solution of silver dihydrogen citrate, as described in United States Patent No. 6,197,814. A summary of the process of making the silver dihydrogen citrate solution is as follows.

A process of making the stock solution of silver dihydrogen citrate is an electrolytic process. The process begins with preparation of an electrolyte solution, which is an aqueous solution comprising citric acid. It has been found that citric acid solutions having citric acid concentrations in the range of about 1% (wt./vol.) to about the solubility limit of citric acid in water (about 60% wt./vol.) are suitable for preparing silver

dihydrogen citrate solutions. In some embodiments, the range is about 5 to about 25 wt% citric acid, with specific values being 10, 15 and 20 wt%. The electrolyte may comprise other ingredients that will not interfere with the electrolytic process.

A pair of silver electrodes are immersed into the electrolyte solution at a suitable spacing to allow ionic current flow between them. A suitable spacing is greater than about 2 mm, for example from about 2 to about 8 mm. An electrolytic potential is applied across the electrodes to create an ionic current flow between the electrodes. While it is possible to use a D.C. power supply to apply the electrolytic potential, it has been found desirable to apply an alternating wave form to the electrodes. A suitable voltage is about 12 to about 50 volts. A suitable alternating waveform applies a pulsed potential having a peak voltage of about 12 to 50 volts to the electrodes, with periodic intermittent phase reversal. The resulting flow of ions through the electrolyte solution results in production of an aqueous solution of silver dihydrogen citrate and citric acid. This solution may be further processed, for example by settling, filtration or both, before being used in the process of making anhydrous silver dihydrogen citrate.

It is possible to perform the foregoing process batch-wise or in a continuous process. It is also possible to recirculate silver dihydrogen citrate solution through the electrolytic cell to increase the final concentration of silver dihydrogen citrate in the solution. The person of skill in the art will recognize that the foregoing steps may be practiced in a number of different variations known in the art. The process is described in fuller detail in United States Patent No. 6,197,814 (incorporated by reference) and in the examples below. The resulting solution is referred to herein as the "stock solution," meaning a water solution of silver dihydrogen citrate and citric acid that has not yet been subjected to freeze-drying.

Having provided the stock solution, the next step is to remove water from the stock solution to produce anhydrous silver dihydrogen citrate. It has been found that freeze-drying results in a superior product. In the freeze-drying process, a frozen solution comprising silver dihydrogen citrate and citric acid solution is subjected to a vacuum, whereby the temperature and pressure of the solution are reduced below the triple point of water. Below the triple point, water will sublime, passing directly from the solid phase into the gaseous phase. In some embodiments, the solution is first frozen, then placed in

a freeze-drying apparatus. The solution may be frozen, for example, by placing it in a cold environment such as a freezer. In other embodiments, the solution is placed in the freeze-drying apparatus and then frozen. As evaporation is an endothermic process, the solution may be frozen by placing it in the freeze-drying apparatus and applying the vacuum. As water evaporates from the solution sample, it draws heat from the sample, thereby lowering the sample's temperature to below the triple point of the sample. In such cases, it is useful to take measures, such as centrifugation, to prevent foaming of the sample. It is also useful to heat the sample during the freeze-drying process in order to compensate for the latent heat deficit of the frozen solution caused by sublimation of water. The person skilled in the art will recognize that heating the frozen sample will generally speed the freeze-drying process, so long as the sample's temperature remains below the triple point of the solution.

Freeze-drying apparatuses are known and available from a number of suppliers; or they may be constructed using procedures known in the art. A freeze-drying apparatus comprises a drying chamber, a condenser to trap water of sublimation, a cooling system that supplies refrigerant to the vacuum chamber and the condenser, a vacuum system to apply a vacuum to the drying chamber. The freeze-drying apparatus may also have a temperature control device that allows one to cool the sample to the freezing point and to heat the sample to speed the sublimation process. The temperature control device will in some cases include refrigeration coils that circulate refrigerant. The temperature control device may also include heating coils or another appropriate device to heat the sample during the freeze-drying process.

The freeze-drying process removes sufficient water to form an anhydrous composition comprising silver dihydrogen citrate and citric acid. The anhydrous silver dihydrogen citrate crystals are translucent-gray in color. In some embodiments, these crystals are ground to a fine powder and stored for an indefinite period of time until they are reconstituted. In some embodiments, the powder is stored in a light-proof container, such as an opaque bottle or in a light-proof box to prevent decomposition by light.

Surprisingly it has been found that a silver dihydrogen citrate composition reconstituted using the same volume of diluent, such as water, as was present in the stock solution possesses the same concentration of free silver ion as was in the stock solution,

without any noticeable degradation, reduction of the redox state of silver, or formation of colloid. As silver ion is the active antimicrobial agent in a silver dihydrogen citrate composition, such reconstituted compositions are expected to possess the same antimicrobial properties as the stock solution. The reconstituted compositions may thus
5 be used in any manner that the stock solutions could have been used.

The term "composition comprising anhydrous silver dihydrogen citrate and citric acid" means a solid composition containing silver dihydrogen citrate and citric acid. Thus, while the composition may contain detectable quantities of water, it does not contain enough water for the composition to be fluid. While the solid composition may
10 be amorphous or crystalline, a crystalline form is especially desirable, as such is formed by the freeze-drying process described, and is considered especially stable. Thus, the invention further provides crystalline compositions comprising anhydrous silver dihydrogen citrate and citric acid.

The invention further provides anhydrous silver dihydrogen citrate compositions
15 that are substantially free of water. The term "substantially free" means that the sample comprises less than the lower detectable limit of water. While it is theoretically possible to remove all water from the sample, in some cases it is sufficient to dry the sample to the point that the amount of remaining water, if any, is below the detection limit for art-recognized detection means. Some detection means that may be mentioned are gas
20 chromatography (GC), nuclear magnetic resonance spectrometry (NMR), mass spectrometry (MS), or combinations of two or more thereof, for example GC-MS.

The invention further provides anhydrous silver dihydrogen citrate containing an amount of water above the detection limit of water by conventional testing processes. In such embodiments, it is preferred that the anhydrous silver dihydrogen citrate comprises
25 less than about 2%, in particular less than about 1%, more particularly less than about 0.5% and even more particularly less than about 0.1% water (wt./wt.%). The person skilled in the art will recognize that such anhydrous compositions may be obtained by performing the freeze-drying process under conditions, such as vacuum, length of time and heating, sufficient to remove the appropriate amount of water from the frozen sample.

30 The invention further provides compositions comprising anhydrous silver dihydrogen citrate and a molar excess of citric acid. The term "molar excess" means an

amount of citric acid such that, for each mole of silver dihydrogen citrate in the composition, there is greater than one mole of citric acid. For example, a 1-60% wt/vol solution of citric acid that has been subjected to electrolysis as previously describe will contain from about 0.00001 to about 1 wt% silver ion. In a particular example, a 10%
5 citric acid solution treated by the electrolysis process described above contains about 0.1% silver ion. The person of skill in the art will recognize that there is a molar excess of citric acid to silver dihydrogen citrate in the solution. In this example, the molar excess is about 15 fold. The person of skill in the art will further recognize that, as citric acid is non-volatile, the solid silver dihydrogen citrate composition produced by the
10 foregoing freeze-drying process will contain a molar excess of citric acid over silver dihydrogen citrate. In some embodiments, this molar excess is from about 5 to about 10,000 fold; in particular about 8 to about 10,000 fold. It is believed that an excess of citric acid over silver dihydrogen citrate of greater than about 5, more particularly greater than about 10, and even more particularly greater than about 12, provides substantial
15 stability of the silver dihydrogen citrate complex in solution. Thus, the invention provides solid silver dihydrogen citrate compositions comprising silver dihydrogen citrate and greater than about 5-, in particular at least about 8-, more particularly at least about 10-, and even more particularly at least about 12-fold molar excess of citric acid over silver dihydrogen citrate.

20 It has surprisingly been found that anhydrous silver dihydrogen citrate compositions are capable of being reconstituted to produce an antimicrobially active composition. It has not been previously reported that an anhydrous silver dihydrogen citrate composition would dissolve in a diluent without disrupting the association between the silver cation and the citrate anion; nor has it been shown that reconstituted solutions
25 have antimicrobial properties.

The invention provides reconstituted silver dihydrogen citrate and citric acid compositions containing from about 0.05 ppm to about 10,000 ppm of silver ion (Ag^+). An effective concentration of silver dihydrogen citrate depends on the microbe against which protection is sought, the substrate to be treated and the existing or potential
30 bioburden on the substrate, and presence or absence of additional antimicrobial actives in the diluent. In some embodiments, an antimicrobially effective concentration of silver

ion ranges from about 0.075 ppm to about 2,500 ppm, especially about 0.1 to about 1,000 ppm.

Thus, the invention provides a method of using an anhydrous silver dihydrogen citrate composition produce a reconstituted silver dihydrogen citrate solution. The method comprises combining the anhydrous composition with a diluent. The diluent is a solvent capable of dissolving the silver dihydrogen citrate and citric acid. In some embodiments, a suitable diluent is an aqueous solvent, such as water or a solution comprising water and an additional water-soluble ingredient. For example the diluent can be water, ethanol or a combination of water and ethanol, such as a 10% solution of ethanol in water.

The invention further provides for reconstituting the anhydrous silver dihydrogen citrate and citric acid compositions with a diluent that is substantially pure water. The term "substantially pure water" means water that is essentially free of contaminants or antimicrobial agents, such as ethanol. Examples of substantially pure water are distilled water, double distilled water, double distilled deionized water, ultrafiltered water and water for injection.

The invention further provides for reconstituting the anhydrous silver dihydrogen citrate and citric acid compositions with tap water, with "fresh water," that is water obtained from rivers, lakes, streams, reservoirs, etc., and with "treated water," that is effluent water that has been treated to remove contaminants. In most normal circumstances, normal tap water will suffice as a diluent, although a purer form of water may be used if necessary.

The invention also provides for reconstituting the anhydrous silver dihydrogen citrate and citric acid compositions with a diluent that is an aqueous solution comprising water and another ingredient. The other ingredient can be a cleaning agent, an antibacterial agent or another type of ingredient. Exemplary cleaning agents include detergents. Exemplary antimicrobials include ethanol. The invention provides diluents comprising water and a member of the group consisting of detergents, alcohols and combinations thereof. Exemplary diluents include aqueous ethanol, aqueous sodium dodecyl sulfate, and aqueous mixtures of ethanol and sodium dodecyl sulfate. For example, the invention provides as a diluent an aqueous solution of about 0.1 to about

10% ethanol and optionally about 0.0001 to about 0.1% detergent. An exemplary diluent comprises about 2% ethanol and greater than about 0.01% detergent in water.

The combination of silver dihydrogen citrate and ethanol produces unexpected synergistic disinfectant and anti-microbial properties. As compared to an aqueous ethanol composition without silver dihydrogen citrate, the invention provides disinfectant and antimicrobial effects at a lower concentration of alcohol than is necessary without the silver dihydrogen citrate.

The invention further provides methods of using anhydrous silver dihydrogen citrate compositions to produce an antimicrobial effect. The methods include combining an anhydrous silver dihydrogen citrate composition with a diluent to produce a reconstituted silver dihydrogen citrate solution. The methods further comprise applying the reconstituted silver dihydrogen citrate and citric acid solution to a substrate. The term "substrate" is generically used herein to mean any surface, article or environment that is in need of antimicrobial treatment.

Various surfaces may be treated with compositions comprising silver dihydrogen citrate, including countertops, floors, glass surfaces, metal surfaces (such as stainless steel, chrome and copper surfaces), tile surfaces, concrete surfaces, vinyl flooring and painted surfaces. The term "surface" is used herein to connote any surface, including interior and exterior surfaces of various objects, including interiors of containers (such as boilers, water tanks, swimming pools, etc.), interiors of pipes, exteriors of household fixtures and appliances, countertops, glass windows and doors. The term "surface" is used herein to distinguish over a whole article. The reconstituted silver dihydrogen citrate and citric acid solutions are applied to the surface by in a conventional manner, such as by pouring, spraying or swabbing the solution onto the surface. The solution are conveniently wiped or rinsed off the surface, or are advantageously left on the surface to dry, thereby providing a long-lasting antimicrobial residue or film on the surface.

The invention further includes methods of treating various articles, such as fabrics, metal articles, plastic articles, natural products and other articles that are often treated with aqueous cleaning solutions. For example, the invention provides for treating food items with a reconstituted silver dihydrogen citrate and citric acid composition. Exemplary food items that are treated in such a manner include vegetables and fruits.

Exemplary vegetables that may be treated in this manner include: roots (such as carrots, beets, radishes); tubers (such as potatoes, turnips, sweet potatoes and yams); bulbs (such as onions, scallions); corms (such as garlic); green leafy vegetables (such as spinach, kale, lettuce and cabbage); cruciferous sprouts (such as broccoli and cauliflower); and legumes (such as beans and peanuts). Exemplary fruits that may be treated in this manner include: squash, melons, apples, peaches, pears, bananas, tomatoes, citrus (such as oranges, grapefruit, tangerines, tangelos, lemons and limes), grapes and olives. The invention provides for spraying or wiping the silver dihydrogen citrate composition onto the food item. The invention alternatively provides for dipping the food item into a the solution.

The invention also embraces methods of treating a variety of articles with reconstituted silver dihydrogen citrate solutions. For example, the solutions may be sprayed or wiped onto a variety of items including mechanical parts used in food service or food process manufacturing. The solutions may also be used to clean toys and other items handled by children and infants. The solutions may further be used to clean fabric items, such as clothing, wash rags, bedding and other fabric items. The invention provides reconstituted silver dihydrogen citrate compositions as soaking compositions for disinfecting cooking utensils, eating utensils and cutlery.

The invention further includes antimicrobially effective silver dihydrogen citrate compositions. The term "antimicrobially effective" means that the composition is effective to halt or reduce the spread of one or more microbes, prevent infestation by one or more microbes, kill one or more microbes, or a combination of these effects. The term microbe includes a bacterium, virus or fungus. Aqueous silver dihydrogen citrate compositions have been shown to be effective against a variety of indicator and pathogenic microbes, including bacteria, such as: *Pseudomonas aeruginosa*, *Salmonella choleraesuis*, *Staphylococcus aureus*, *Propionibacterium acnes*, *Escherichia coli*, *Listeria monocytogenes* and *Enterococcus faecium*; viruses such as: Human immunodeficiency virus 1 (HIV 1), Herpes simplex virus 2 (HSV 2), Influenza A, Rhinovirus and Poliovirus type 2; and fungi such as *Trichophyton mentagrophytes*.

The invention may be more fully appreciated with reference to the following illustrative and non-limiting examples. Other embodiments of the invention may be practiced within the scope of the present invention.

EXAMPLES

Example 1 – Preparation of Stock Silver Dihydrogen Citrate Solution

Water was introduced into a reverse osmosis unit, passing through a semi-permeable membrane to remove impurities and producing deionized water. Anhydrous
5 99% pure citric acid was mixed with the water to produce 200 gallons of a 20 % (wt/vol)
(796 g citric acid per gallon water) solution. The 200 gallons of 20% citric acid were
directed into an ion chamber containing having positive and negative electrodes, each
consisting of 200 troy ounces of 999 fine silver. The positive and negative electrodes
were spaced at least 2.0 mm apart, allowing the citric acid solution to pass between the
10 two electrodes. An ion generation controller (IGC) power supply including a positive and
a negative conductor was attached to the positive and negative electrodes. The IGC
applied a current of 5 amps at 17 volts, pulsed every 9 seconds, with a polarity change at
1 minute intervals. Throughout the process, the electrode gap was adjusted in order to
maintain the 5 amp-17 volt output. The electric current flow caused an ion current to
15 flow between the positive and negative electrodes, producing free silver ions within the
diluted citric acid solution. The silver ions reacted with the citric acid in the citric acid
solution to produce the silver dihydrogen citrate solution. The 20% citric acid solution
was recirculated through the ion chamber at 50 gallons per minute for 144 hours until the
desired silver ion concentration was obtained. The silver dihydrogen citrate solution was
20 then allowed to sit in order to allow any solids formed during the procedure to precipitate.
The resulting product was a silver dihydrogen citrate solution having a silver ion
concentration of 2400 ppm. Hereafter, this solution is referred to as a stock solution.
The stock solution can be used immediately per the following examples or stored for later
use.

25 It should be understood by those skilled in the art that numerous variations in the
size and/or spacing of the electrodes and numerous variations in the peak voltage and
numerous variations in the timing sequence of the intermittent voltage polarity can readily
be used to obtain the silver dihydrogen citrate for use in the invention.

Example 2 – Preparation of Anhydrous Silver Dihydrogen Citrate

30 A 1,000 ml sample of the stock solution, prepared as in foregoing Example 1, was
obtained from SSA Batch No. 04.06.03 and was confirmed to have a concentration of

ionic silver of 2400 ppm using an Orion™ 290A processor with an ion specific electrode (ISE). The processor was calibrated using Orion™ standard solutions.

The 1,000 ml stock solution was decanted into two 2.5" x 8" x 15" Pyrex™ heat resistant glass trays. The liquid stock solution level was approximately 1.5" in each tray.

5 The trays were then placed into a small commercial-grade freeze-drying unit, which consisted of a drying chamber with temperature control shelves, a condenser to trap water removed from the product, a cooling system to supply refrigerant to the temperature control shelves and condenser, a vacuum system to reduce the pressure in the chamber and condenser to facilitate the drying process. Cooling and vacuum pressure
10 settings were kept within the standard ranges associated with freeze-drying aqueous solutions > 50% water matrix.

After 99% of the water was removed from the product, the large, coarse crystals were collected from the trays and ground into a fine crystalline powder, weighed and stored at an ambient room temperature (74°F) for one week. Total crystal weight was
15 206.12 g.

Example 3 – Reconstitution of Silver Dihydrogen Citrate Solution

The crystalline powder from Example 2 was then reconstituted using approximately 794 ml of pharmaceutical-grade pure water in a 1,000 ml Pyrex glass flask. The solution was then agitated via magnetic stirrer for 30 minutes and put into a
20 light-proof storage cabinet for 24 hours.

The concentration of silver ion in the reconstituted solution was then measured using the Orion 290A processor and ISE described in Example 2. Measurements were obtained at 24, 48 and 72 hours and at intervals of 1 week for a total of 4 weeks. The 24 hour reading and the week 4 readings were 2398 ppm and 2407 ppm, respectively. These
25 concentrations are essentially the same as the 2400 ppm stock solution (within the % error operational specifications for the Orion 290A™ processor).

Example 4 – Antimicrobial Efficacy of Reconstituted Silver Dihydrogen Citrate Solutions.

In order to demonstrate the efficacy of silver dihydrogen citrate compositions as
30 preservatives, stock solutions of silver dihydrogen citrate, as described in Example 1,

were subjected to Preservative Challenge Tests. The Preservative Challenge Tests were performed according to the European Pharmacopoeia test method 4.04/5.01.03.00 for Category 2 products (topically used products made with aqueous bases or vehicles, nonsterile nasal products, emulsions including those applied to mucous membranes.

- 5 Bacterial test organisms and yeasts were cultivated on Casein Soymeal peptone agar and fungal test organisms on Sabouraud 4% glucose agar for 18-24 hours at 35°C (bacteria), 48 hours at 25°C (Candida) or 1 week at 25°C (Aspergillus).

- After incubation, the bacterial and yeasts were harvested by washing off the surface of the agar plates with 0.9% sodium chloride. Aspergillus was harvested by
10 washing off the agar plate surface with 0.9% sodium chloride /0.01% Tween 80.

 The suspension of test microorganisms were diluted with 0.9% sodium chloride to the final test organism suspensions with a density of 10^8 colony forming units.

- Per test organisms, 20g of the test product were weight in glass jars (250 ml jars with screw cups from Schott/Germany) and contaminated with 0.2 ml of the test organism
15 suspension. The microorganisms were carefully distributed in the test product by stirring with a glass spatula.

 The so contaminated test products were stored at 20-25°C in the dark.

 Samples of 1 g material were taken immediately after contamination of the test products and 2 days, 7 days, 14 days and 28 days after contamination.

- 20 The samples were diluted in 0.9% sodium chloride and 0.1 ml aliquots of the dilutions were spread on agar plates by means of Drigalsky spatula. An adequate inactivator (neutralizer) of the specific antimicrobial was incorporated in the diluent used for preparation of the product dilutions and in the agar plates used for assessment of the total number of viable cells.

- 25 The agar plates were incubated for 24 hours at 35°C (bacteria and yeasts) or 3 days at 25°C (Aspergillus) and the grown colonies were counted after the incubation phase. The colonies were counted and the number of viable cells (colony forming units) per g test product was calculated. The log reduction of the microorganisms in the product was then calculated (see tables with results of Preservation Challenge Tests below).

Test strains:

Pseudomonas aeruginosa ATCC 9027; NCIMB 8626; CIP 82.118

Staphylococcus aureus ATCC 6538; NCTC 10788; NCIMB 9518; CIP 4.83

Candida albicans ATCC 10231; NCPF 3179; IP 48.72

5 *Aspergillus niger* ATCC 16404; IMI 149007; IP 1431.83

Silver dihydrogen citrate was tested in a variety of formulations for its antimicrobial effects. The following Tables 1-3 show the results of these tests:

Table 1**Preservative Challenge Test / Deodorant Emulsion**

Test organisms	Staph. aureus	E. coli	Ps. aeruginosa	C. albicans	A. niger
O/W PK03-260-01 (Placebo)					
2 days after contamination	< 100	< 100	< 100	6.0 x 10E4	1.8 x 10E5
7 days after contamination	< 100	< 100	< 100	1.6 x 10E4	3.0 x 10E5
14 days after contamination	< 100	< 100	< 100	8.8 x 10E3	2.8 x 10E5
28 days after contamination	< 100	< 100	< 100	1.4 x 10E3	n.d.
O/W PK03-260-01 (0.1% Axenohl)					
2 days after contamination	< 100	< 100	< 100	< 100	1.6 x 10E5
7 days after contamination	< 100	< 100	< 100	< 100	2.6 x 10E5
14 days after contamination	< 100	< 100	< 100	< 100	1.2 x 10E5
28 days after contamination	< 100	< 100	< 100	< 100	n.d.
O/W PK03-260-01 (0.3% Axenohl)					
2 days after contamination	< 100	< 100	< 100	< 100	1.4 x 10E5
7 days after contamination	< 100	< 100	< 100	< 100	1.2 x 10E5
14 days after contamination	< 100	< 100	< 100	< 100	1.0 x 10E5
28 days after contamination	< 100	< 100	< 100	< 100	n.d.

Table 2**Preservative Challenge Test / Deodorant Emulsion**

Test organisms	Staph. aureus	E. coli	Ps. aeruginosa	C. albicans	A. niger
PK03-262-01 (Placebo)					
2 days after contamination	2.0 x 10E4	1.0 x 10E5	1.0 x 10E6	1.0 x 10E6	2.0 x 10E5
7 days after contamination	< 100	3.1 x 10E3	4.4 x 105	6.2 x 10E5	3.0 x 10E5
14 days after contamination	< 100	1.0 x 10E2	1.1 x 10E8	1.3 x 10E6	3.5 x 10E5
28 days after contamination	< 100	1.2 x 10E2	6.0 x 10E7	1.1 x 10E6	n.d.
PK03-262-01 (0.1%AxenohI)					
2 days after contamination	5.0 x 10E2	4.0 x 10E2	3.0 x 10E2	4.0 x 10E3	1.4 x 10E5
7 days after contamination	< 100	2.0 x 10E2	< 100	3.4 x 10E4	1.8 x 10E5
14 days after contamination	< 100	< 100	< 100	1.1 x 10E5	2.0 x 10E5
28 days after contamination	< 100	< 100	< 100	4.0 x 10E5	n.d.
PK03-262-01 (0.3%AxenohI)					
2 days after contamination	3.0 x 10E2	4.0 x 10E3	7.0 x 10E2	4.0 x 10E3	1.8 x 10E5
7 days after contamination	< 100	2.0 x 10E2	< 100	2.4 x 10E3	2.4 x 10E5
14 days after contamination	< 100	100	< 100	1.2 x 10E3	2.4 x 10E5
28 days after contamination	< 100	< 100	< 100	2.0 x 10E2	n.d.

Table 3**Preservative Challenge Test / Shower Gel**

Test organisms	Staph. aureus	E. coli	Ps. aeruginosa	C. albicans	A. niger
FB02-060-03 (Placebo)					
2 days after contamination	4.0 x 10E2	2.2 x 10E5	in progress	2.2 x 10E5	2.4 x 10E5
7 days after contamination	< 100	1.1 x 10E5	in progress	2.1 x 10E5	3.8 x 10E5
14 days after contamination	< 100	8.2 x 10E4	in progress	9.0 x 10E4	3.8 x 10E5
28 days after contamination	n.d.	n.d.	in progress	n.d.	n.d.
Body Shampoo (0.1% Axenohl)					
2 days after contamination	< 100	1.8 x 10E4	< 100	< 100	2.0 x 10E5
7 days after contamination	< 100	< 100	< 100	< 100	3.0 x 10E5
14 days after contamination	< 100	< 100	< 100	< 100	3.2 x 10E5
28 days after contamination	< 100	< 100	< 100	< 100	n.d.
Body Shampoo (0.3% Axenohl)					
2 days after contamination	< 100	8.0 x 10E3	< 100	< 100	1.8 x 10E5
7 days after contamination	< 100	< 100	< 100	< 100	2.4 x 10E5
14 days after contamination	< 100	< 100	< 100	< 100	2.4 x 10E5
28 days after contamination	< 100	< 100	< 100	< 100	n.d.

The invention may be more fully appreciated with reference to the following illustrative and non-limiting examples. Other embodiments of the invention may be practiced within the scope of the invention.

What is claimed is:

1. A composition comprising anhydrous silver dihydrogen citrate and citric acid.
2. The anhydrous composition of claim 1, wherein the composition is substantially free of water.
- 5 3. The anhydrous composition of claim 1, wherein the composition contains less than about 0.5% water.
4. The anhydrous composition of claim 1, wherein the composition is crystalline.
5. The anhydrous composition of claim 1, wherein the composition comprises a molar excess of citric acid over silver dihydrogen citrate.
- 10 6. The anhydrous composition of claim 1, wherein the composition comprises at least a 5 fold molar excess of citric acid over silver dihydrogen citrate.
7. A process of making a composition comprising an anhydrous silver dihydrogen citrate and citric acid, comprising the steps of:
 - 15 (a) providing a stock solution comprising silver dihydrogen citrate, citric acid and water;
 - (b) freezing the stock solution to form a frozen solution; and
 - (c) freeze-drying the frozen solution to form the anhydrous composition.
8. The process of claim 7, wherein the citric acid is a molar excess of citric acid over silver dihydrogen citrate.
- 20 9. The process of claim 7, wherein step (c) further comprises heating the frozen solution.
10. A method of using an anhydrous silver dihydrogen citrate and citric acid composition, comprising combining the anhydrous silver dihydrogen citrate composition with an amount of aqueous diluent sufficient to produce a reconstituted antimicrobially active silver dihydrogen citrate composition.
- 25 11. The method of claim 10, wherein the aqueous diluent is substantially pure water.

12. The method of claim 10, wherein the aqueous diluent comprises water and at least one member of the group consisting of alcohols and detergents.
13. A method of using an anhydrous silver dihydrogen citrate and citric acid composition, comprising combining the anhydrous silver dihydrogen citrate composition with sufficient aqueous diluent to produce an antimicrobially effective silver dihydrogen citrate solution; and applying the antimicrobially effective silver dihydrogen citrate solution to a substrate, whereby an antimicrobial effect is achieved.
14. The method of claim 13, wherein the substrate is a solid surface, a food article or a non-food article.
15. The method of claim 14, wherein the solid surface is the interior of a pipe, a holding tank, a swimming pool, a spa, a cooling system or a cooling tower.
16. The method of claim 13, wherein the antimicrobial effect is biocidal or biofilm control.
17. The method of claim 13, wherein the antimicrobial effect is food preservation.